FDA 510(k) Premarket Notification

K952860 THE NINER NITINOL GUIDEWIRE

CLEARED

Dec 13, 1995
174 days to decision

K952860 · Product code: **DQX** · Cardiovascular
Source: https://www.510kdatabase.net/k952860/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Wire, Guide, Catheter (DQX)
Date received	Jun 22, 1995
Decision date	Dec 13, 1995
Days to decision	174 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Microvena Corp.**
Location	Findley, MN, US
Contact	JODI LOCHER
510(k) history	18 submissions · 18 cleared · 1990-1999